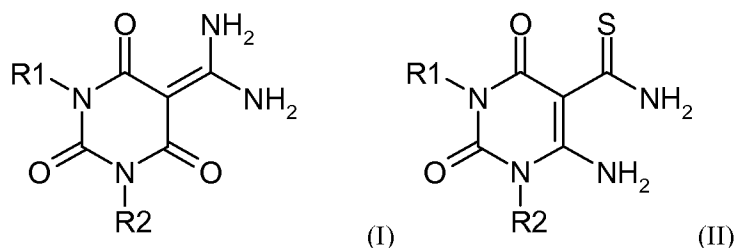


ABSTRACT

This invention relates to compounds of formula (I) or (II)



wherein, in either formula, independently;

R¹ and R² are the same or are different and are C₁₋₈ alkyl, C₂₋₈ alkylene, C₃₋₈ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₃₋₆ cycloalkylaryl, or heterocycloaryl; wherein said alkyl, alkylene, cycloalkyl, aryl, heteroaryl, heterocyclyl, cycloalkylaryl, or heterocycloaryl are unsubstituted or substituted by one or more groups selected from the group consisting of halogen, C₁₋₈ alkyl, C₁₋₈alkoxy, C₁₋₈thioalkoxy, cycloalkyl, aryl, heteroaryl, heterocycloalkyl, CF₃, SCF₃, NHC(O)_nR⁵, S(O)_mR⁵, S(O)₂NR⁵R⁶, C(S)NR⁵R⁶, CONR⁵R⁶, C(O)_nR⁵;

n is 0, 1 or 2;

m is 0, 1 or 2;

R⁵ is hydrogen, alkyl, aryl, alkylaryl, heterocycloalkyl, or heteroaryl and is unsubstituted or substituted by one or more groups selected from the group consisting of alkyl, C₁₋₈alkoxy, aryl, heteroaryl, halogen, NO₂, CN, N₃, SCF₃, and CF₃;

R⁶ is hydrogen, alkyl, aryl, alkylaryl, heterocycloalkyl, or heteroaryl and is unsubstituted or substituted by one or more groups selected from the group consisting of alkyl, C₁₋₈alkoxy, aryl, heteroaryl, halogen, NO₂, CN, N₃, SCF₃, and CF₃, or when R¹ and/or R² contains S(O)₂NR⁵R⁶, CONR⁵R⁶, or C(S)NR⁵R⁶, then R⁵R⁶ together with the nitrogen may form a heterocyclic ring; or

a pharmaceutically acceptable salt or solvate thereof.